Comprehensive plaque assessment by coronary CT angiography

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Abstract | Most acute coronary syndromes are caused by sudden luminal thrombosis due to atherosclerotic plaque rupture or erosion. Preventing such an event seems to be the only effective strategy to reduce mortality and morbidity of coronary heart disease. Coronary lesions prone to rupture have a distinct morphology compared with stable plaques, and provide a unique opportunity for noninvasive imaging to identify vulnerable plaques before they lead to clinical events. The submillimeter spatial resolution and excellent image quality of modern computed tomography (CT) scanners allow coronary atherosclerotic lesions to be detected, characterized, and quantified. Large plaque volume, low CT attenuation, napkin-ring sign, positive remodelling, and spotty calcification are all associated with a high risk of acute cardiovascular events in patients. Computation fluid dynamics allow the calculation of lesion-specific endothelial shear stress and fractional flow reserve, which add functional information to plaque assessment using CT. The combination of morphologic and functional characteristics of coronary plaques might enable noninvasive detection of vulnerable plaques in the future.

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Introduction
Cardiovascular diseases are the number one cause of death globally. The number of people who die from cardiovascular diseases, mainly from coronary heart disease and stroke, will increase to reach 23.3 million by 2030 from an estimated 17.3 million deaths in 2008. Cardiovascular diseases are projected to remain the single leading cause of death by 2030 globally. In 2014, an estimated 1.1 million people in the USA will have a major adverse coronary event, which will lead to death in ~34% of those individuals, despite many effective treatment strategies existing beyond the coronary lumen and stenosis is of utmost importance as we develop personalized medical care to guide prevention, medical therapy, and coronary interventions. The ability to interpret CCTA images beyond the coronary atherosclerotic plaque, not just the coronary lumen. CCTA provides information regarding the coronary tree and atherosclerotic plaques beyond simple luminal narrowing and plaque type defined by calcium content. These novel applications will improve image-guided prevention, medical therapy, and coronary interventions. The ability to interpret CCTA images beyond the coronary lumen and stenosis is of utmost importance as we develop personalized medical care to enable therapeutic interventions stratified on the basis of plaque characteristics.

In this Review, we describe the morphological and functional features of vulnerable plaques, as potential

Competing interests
The authors declare no competing interests
REVIEWS

**Key points**
- Most acute coronary events result from sudden luminal thrombosis due to rupture of an atherosclerotic plaque
- Modern computed tomography (CT) scanners enable robust coronary plaque characterization and quantification
- Large plaque volume, low CT attenuation, napkin-ring sign, positive remodelling, and spotty calcification are all associated with plaques vulnerable to rupture
- Computational fluid dynamic simulation enables plaque-specific endothelial shear stress and fractional flow reserve assessment, and thus permits functional characterization of plaques
- Coupling individual plaque morphology with plaque-specific functional data will enable new noninvasive detection of vulnerable plaques with CT

**Plaque morphology**

Histological investigations have revealed three distinct features of plaques associated with acute coronary events: rupture; erosion; and calcified nodule. Two-thirds of luminal thrombi in acute events result from ruptured atherosclerotic lesions characterized by a necrotic core covered by a thin layer of fibrous cap (Figure 1b). Plaques vulnerable to rupture might have the same morphological characteristics as ruptured plaques, but with an intact thin fibrous cap. These lesions—termed thin-cap fibroatheroma (TCFA), with a cap thickness of <65 μm—are considered to be the precursor lesions of plaque rupture. The limited spatial resolution of current CT scanners (~400 μm) precludes the morphometric analysis of fibrous cap by CCTA. Histopathological investigations suggest that plaques prone to rupture are enlarged in all three spatial dimensions. In TCFAs the necrotic core length is ~2–17 mm (mean 8 mm) and the area of the necrotic core in 80% of cases is >1.0 mm². These dimensions are over the plaque detection threshold (>1 mm plaque thickness) for CCTA. Moreover, the majority of TCFAs occur in the proximal portions of the main coronary arteries, where vessel diameter is largest, and CCTA has the highest image quality and accuracy for the plaque detection.

**Coronary plaque burden**

Large plaque volume was associated with the diagnosis of ACS in cross-sectional studies, and quantification of noncalcified plaques (NCP) can improve risk stratification and improve the prognostic value of CCTA to predict future cardiovascular events. The multicentre Providing Regional Observations to Study Predictors of Events in the Coronary Tree (PROSPECT) trial is the first and largest natural-history study of coronary plaques using invasive angiography and intravascular ultrasound (IVUS) to identify plaques vulnerable to rupture on a per-lesion basis. The prospective study included 697 patients with ACS in whom 3-vessel grey-scale IVUS and IVUS with radiofrequency backscatter analysis (known as virtual histology IVUS [VH-IVUS]) were performed to characterize nonculprit (that is unruptured) lesions. After a median of 3.4 years follow-up, the strongest predictor of future events was the IVUS-derived plaque burden of ≥70% (HR 5.03; 95% CI 2.51–10.11; P<0.001). CT data sets can provide submillimeter isotropic spatial resolution, and the possibility of CT attenuation-based tissue characterization enable the quantification of total coronary plaque burden and individual plaque components, which is similar to the results obtained with IVUS. Automated software tools are now available for plaque quantification and characterization (Figure 2).

Automated quantification of plaques is desirable to improve the reproducibility, accuracy and efficiency of CCTA plaque analysis. The reproducibility of automated 3D quantification software for plaque burden was

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**Figure 1** | The morphology and functional characteristics of stable and vulnerable plaques. **a** | Stable fibrocalcific lesion with calcification and small lipid pools. The plaque leads to mild narrowing of the lumen; however, there is no ischaemia after the lesion (FFR >0.8; green). ESS near the plaque is in the normal physiological range indicating undisturbed flow. **b** | Rupture prone vulnerable plaque with a large lipid-rich necrotic core, thin fibrous cap, neovascularization, spotty calcium and presence of inflammatory cells. Despite the positively remodelled vessel wall at the site of the plaque, the lesion causes severe luminal narrowing and ischaemia (FFR <0.8; red). The downstream plaque region with low and oscillatory ESS promotes plaque growth, whereas the upstream low ESS at the shoulder regions is more inflamed (indicated by presence of macrophages), which might lead to plaque destabilization. High ESS at the most stenotic part can trigger plaque rupture. Abbreviations: ESS, endothelial shear stress; FFR, fractional flow reserve. Part b modified with permission from Nature Publishing Group © Slager, C. J. et al. The role of shear stress in the destabilization of vulnerable plaques and related therapeutic implications. Nat. Clin. Pract. Cardiovasc. Med. 2, 456–464 (2005).
demonstrated to be excellent, with an intraclass correlation (ICC) value of 0.88 (95% CI 0.74–0.95); excellent agreement was defined as an ICC coefficient of >0.8.\textsuperscript{30} The accuracy of automated coronary plaque quantification by CCTA was successfully validated against greyscale IVUS and VH-IVUS.\textsuperscript{30–32} However, automated plaque quantification software tools have poor interplatform reproducibility; the same software should, therefore, be used for serial or comparative assessments.\textsuperscript{33}

The culprit lesion (that is ruptured) and vulnerable plaques evaluated by histology or invasive imaging techniques tend to be large in size, leading to the hypothesis that CCTA quantification might incrementally improve risk stratification of patients over conventional CCTA reading.\textsuperscript{13} A cross-sectional clinical investigation demonstrated that the culprit plaques in patients with ACS have larger volume than stable lesions in patients with stable angina pectoris (SAP; 193 mm$^3$ versus 104 mm$^3$; $P=0.001$).\textsuperscript{34} In patients with unstable angina, quantitative CCTA revealed that plaques with morphological features of plaque disruption (such as intraplaque contrast dye penetration) had a larger volume compared with plaques that had no signs of disruption (313 ± 356 mm$^3$ versus 118 ± 93 mm$^3$; $P<0.0001$). These lesions also contained more low CT attenuation components characteristic to lipid-rich plaques (99 ± 161 mm$^3$ versus 19 ± 18 mm$^3$;

Figure 2 | Example of plaque characterization and quantification using a dedicated automated software tool and CCTA data set. a | Segmented whole coronary tree. The LAD is indicated in blue. The coronary centrelines and the aorta are indicated in green. Red box indicates plaque of interest. b | Curved multiplanar reconstruction of the LAD. Dotted lines indicate a partially calcified, positively remodelled plaque in the LMS bifurcation. c | The LMS plaque cross-section from panel b. d | The LMS plaque cross-section with colour overlay derived with adaptive threshold setting. The lipid rich (low CT attenuation) plaque components are shown in red. Fibro-fatty tissue is shown in light-green. Fibrous tissue is shown in dark-green. Calcium is shown in white. e | Volumetric assessment of the lipid rich plaque core is shown in red, and the core’s spatial relation to the lumen (grey mesh) and calcium (white). f | The graph illustrates the areas of different plaque components. The colour scheme is identical to panel d. Abbreviations: Ca, calcium; CCTA, coronary computed tomography angiography; LAD, left anterior descending artery; LCx, left circumflex coronary artery; LMS, left main stem.
In patients with acute chest pain and obstructive coronary lesions, the total volume of plaques leading to stenosis was not significantly different between those individuals with and those without ACS (212 ± 71 mm³; P = 0.24). Interestingly, the volume of NCP with low CT attenuation density, (<90 Hounsfield units [HU], a measure of CT attenuation) was significantly larger in patients with ACS compared with patients who did not have ACS (91 ± 49 mm³; P = 0.03).16

Longitudinal clinical investigations indicate a strong prognostic value of CCTA derived plaque volume for future coronary events. In a retrospective study of 1,059 patients with stable chest pain, the coronary plaque volume was larger in those patients who developed ACS compared with patients who did not during a follow-up period of 27 ± 10 months (134.9 ± 14.1 mm³ versus 57.8 ± 5.7 mm³; P < 0.001). Nevertheless, the authors of a study published in 2013 elegantly demonstrated that semiautomatic plaque quantification—whereby plaques are manually identified before automatic segmentation, characterization, and quantification, with optional manual corrections—provided additional prognostic value for ACS over both clinical risk factors and traditional CT reading (including calcium score, segment stenosis score, lesion severity, and number of segments with NCP).17 The patients who developed ACS had a higher total plaque volume (median 94 ± 29 mm³; P < 0.001) and total NCP volume (28 ± 4 mm³; P < 0.001) at baseline compared with those individuals who did not develop ACS.17 The volume of nonobstructive NCP measured by CCTA was a strong predictor of future coronary events in a prospective study of 312 patients with non-ST-segment elevation myocardial infarction who underwent CCTA before invasive coronary angiography.18 In total, 23 patients had a coronary event after a median follow-up of 16 months and the total volume of nonobstructive NCP was independently associated with the events with an HR of 1.18 per 100 mm³ plaque volume increase. Interestingly, neither Agatston score (a measure of calcium content in a CT scan), nor calcified plaque volume were associated with an increased risk of coronary event. A subset of patients (n = 32) underwent CCTA in the PROSPECT study.24,38 The authors of this exploratory substudy observed a higher total atheroma volume at baseline in patients with subsequent cardiac events during the mean 39 months follow-up (970 ± 811 mm³; P < 0.01).18

However, despite these promising results, further software improvements are warranted to maximize accuracy, reproducibility, and time-efficiency before automated plaque burden quantification is implemented in the clinic. Moreover, industry standards should be developed to enable reproducible plaque assessments with CCTA regardless of the software tool used.31

Low CT attenuation plaques
Lesions leading to ACS often have a large necrotic lipid-rich core; therefore, the CT differentiation between plaques containing lipid-rich material and plaques with predominantly fibrous components is desirable for prediction of ACS.19 Traditionally, CCTA classifies plaques according to the presence or absence of calcified components, thereby differentiating between calcified, partially-calcified (mixed), and NCPs. The differentiation between calcified plaque components and NCPs was feasible even with early multidetector CT technology (such as 4-slice CTs used in the late 1990s). However, the classification of NCPs into lipid-rich and fibrous lesions on the basis of CT attenuation values (measured by HU) remains challenging.

Some investigators have correlated CCTA plaque assessment with the clinical reference standard IVUS, and report low CT attenuation on average for lipid-rich plaques.43 NCPs with high CT attenuation correlated with fibrous tissue and those with low densities correlated with necrotic core and fibrofatty tissue as assessed by VH-IVUS. In histogram analysis of the intraplaque pixel CT numbers, lipid-rich plaques have a higher percentage of pixels with low HU values compared with plaques of predominantly fibrous components.44 This observation was validated in an ex vivo study that showed that the relative area (area >25%) of intraplaque pixels with <60 HU could accurately detect lipid-rich atherosclerotic lesions (sensitivity, 73%; specificity, 71%). Moreover, low CT numbers were measured in TCFAs identified by optical coherence tomography (OCT; the standard clinical reference for fibrous cap thickness measurements and necrotic lipid-rich core detection) compared with stable lesions (35–45 HU versus 62–79 HU; P < 0.001). However, the variability of CT values within plaque types is wide. Despite the differences in mean densities between fibrous plaques and lipid-rich plaques, almost all investigators have reported a substantial overlap of densities, which prevented the reliable subclassification of NCPs. Furthermore, CT measurements of coronary plaques are influenced by several factors, such as the concentration of adjacent intraluminal iodinated contrast agent, plaque size, image noise, tube voltage, slice thickness, and the reconstruction filter. The reliable differentiation between lipid-rich and fibrous lesions made solely on the basis of CT attenuation is, therefore, not yet feasible. New automated plaque quantification software tools, with scan-specific adaptive attenuation threshold settings, can potentially overcome some of these limitations and might improve CT number-based plaque component quantification (Figure 2).

Despite the challenges associated with CT number-based plaque characterization, low CT attenuation seems to be a consistent feature of lipid-rich plaques. Low-density plaques, defined by <30 HU average attenuation, were more often seen in patients with ACS than in those individuals with SAP (79% versus 9%; P < 0.0001). The same investigators compared the characteristics of ruptured fibrous cap culprit lesions in patients with ACS with the intact fibrous cap plaques of patients with SAP. Again, the low plaque attenuation was defined as <30 HU, and 88% of ruptured plaques had a low CT attenuation, compared with 18% of the stable lesions (P < 0.001). Similarly, other investigators have also reported lower
Napkin-ring sign

Histopathological analysis of culprit and nonculprit TCFAs with similar stenosis revealed that only fibrous cap thickness (OR 0.35; P < 0.05), and necrotic core size (OR 2.0; P < 0.02) are independent predictors of plaque rupture.11 A further analysis of the hierarchical importance of plaque features that are accessible by non-invasive imaging revealed that the size of necrotic core and the presence of macrophage infiltration are the two best discriminators between ruptured plaques or TCFAs and stable lesions.14,15 Furthermore, a large necrotic core cross-sectional area (>3.5 mm²) can differentiate a ruptured plaque or TCFAs from a stable plaque.15 In 80% of plaques vulnerable to rupture, the area of necrotic core is >1.0 mm².19 These dimensions are over the detection threshold of CCTA and, therefore, enable noninvasive risk stratification of individual coronary plaques according to the presence or absence of large necrotic core. A plaque cross-section with low CT attenuation in the central part of the lesion might be indicative of the presence of a large lipid-rich necrotic core.

A ring-like CT attenuation pattern of NCP was observed in ACS culprit plaques and in plaques associated with slow-flow and no re-flow phenomenon during percutaneous coronary intervention.34,46,47,57–59 The term napkin-ring sign (NRS) is used to describe this specific plaque attenuation pattern.60 The NRS is a qualitative plaque feature and can be defined in a noncalcified plaque cross-section by the presence of two features: a central area of low CT attenuation that is apparently in contact with the lumen; and a ring-like higher attenuation plaque tissue surrounding this central area (Figure 3).50,61 Interestingly, NRS was present in both native (that is non-contrast-enhanced) and contrast-enhanced ex vivo CT images, suggesting that the feature is the result of differences in CT attenuation between the large necrotic core (a central low CT attenuation) and fibrous plaque tissue (ring-like higher attenuation).60–62 However, in vivo, some additional factors (such as the vasa vasorum) might influence the development of NRS.57 The area of necrotic core can be over twice the size in NRS plaques compared with non-NRS plaques (median 1.10 mm² versus 0.46 mm²; P = 0.05).62 These values correlate with other histopathological observations that demonstrate the area of necrotic core in vulnerable plaques is >1.0 mm² in the majority (~80%) of cases.59 In CT, the specificity of an NRS to identify an advanced coronary plaque and TCFAs is excellent (98.9% and 94.1%, respectively).63 In a detailed plaque attenuation pattern analysis, a pattern-based plaque classification scheme has been proposed that classifies noncalcified plaques into homogeneous and heterogeneous, and stratifies heterogeneous plaques into NRS and non-NRS lesions (Figure 4).64,65 The diagnostic performance of pattern classification to identify advanced lesions was significantly better than conventional plaque classification made on the basis of calcified content (area under the receiver operating characteristics curve: 0.761 versus 0.678; P < 0.001).41 In clinical investigations the NRS had 96–100% specificity for the identification of TCFAs or culprit ACS lesions.34,66 The NRS was more frequent in TCFAs compared with non-TCFA plaques defined by OCT in two different clinical studies (TCFA 44–65% versus non-TCFA 4–16%; P < 0.0001).66–68 NRS also independently predicted future ACS events (independently of positive remodelling and low attenuation) in a prospective study of 895 patients with a mean follow-up of 2.3 years (HR 5.6; P < 0.001).64

The NRS seems to be a specific CT feature of plaques with a large necrotic core. However, the sensitivity of NRS is relatively low. A more-detailed analysis of different plaque attenuation patterns might provide means for the development of plaque classification schemes with improved diagnostic performance and identify vulnerable coronary plaques.

Positive remodelling

Rupture-prone plaques might not lead to significant luminal narrowing, owing to the effect of positive remodelling.7 Positive remodelling describes the compensatory enlargement of the vessel wall that occurs at the site of the atherosclerotic lesion as the plaque size increases, resulting in the preservation of luminal area.65 In histopathology studies, positive remodelling is associated with the abundance of macrophages and increased necrotic core.66 CCTA can measure the outer vessel wall...
and lumen dimension. The remodelling index is calculated as the vessel cross-sectional area at the site of maximal stenosis divided by the average of proximal and distal reference segments’ cross-sectional areas. A remodelling index threshold of ≥1.1 was suggested for the definition of positive remodelling visualized by CCTA, whereas some authors use ≥1.05 or >1.0 as the cut-off point on the basis of IVUS studies. Automated software now permits the easy quantification of the remodelling index. The remodelling index assessed by CCTA correlates well with IVUS measurements; however, CCTA has a trend towards overestimation of remodelling index (95% CI of the mean difference 0.01–0.08; P = 0.005). Consistent with histopathological data, lesions with positive remodelling on CCTA have a higher plaque burden, a larger amount of necrotic core and a higher prevalence of TCFA assessed by VH-IVUS when compared to lesions without positive remodelling. Furthermore, in two correlative studies comparing CCTA with OCT, the CT-derived remodelling index was higher in TCFA compared with non-TCFA lesions classified by OCT (1.14 versus 1.02, P < 0.0001; and 1.14 versus 0.95, P < 0.0001). A remodelling index threshold of 1.08, which had the best diagnostic performance to identify TCFA, has been suggested by some investigators. In a study of 38 patients with ACS and 33 patients with SAP, positive remodelling was strongly associated with culprit plaques in ACS (87%), but not SAP (12%; P < 0.0001), and had the best diagnostic performance among other high-risk CT plaque features (low attenuation and spotty calcification) to identify the culprit lesions (sensitivity 87%; specificity 88%). Several other cross-sectional CCTA studies have also found a higher remodelling index in patients with ACS compared with patients with SAP (1.14–1.6 versus 0.9–1.2; P = 0.001–0.04). Positive plaque remodelling and/or low plaque attenuation was an independent predictor of ACS in a clinical study with 27 ± 10 months follow-up (HR 22.8; 95% CI 6.9–75.2; P < 0.001). Among patients with one of these high-risk CT features, one in five will have an adverse coronary event within 1–3 years, a similar rate to those with a three-feature positive plaque determined by VH-IVUS in the PROSPECT trial. The remodelling index can be reliably measured by CCTA. However, a more-conservative remodelling index threshold of 1.1 is preferred in the assessment of CCTA images.

Spotty calcium in plaques

Calcification is an ever-present feature of advanced coronary atherosclerosis. Coronary calcification assessed by CT is highly associated with plaque burden and related to poor clinical prognosis. However, the effect of calcification on plaque instability is controversial. Although most acute plaque ruptures in individuals with sudden cardiac death contain some calcification under histopathology, approximately two-thirds have only microcalcification, which is not detectable by CT. In a serial IVUS study, plaques with heavy calcification are clinically quiescent, whereas spotty (small) calcification was associated with accelerated disease progression in patients with SAP. Furthermore, the presence of spotty calcification was related to culprit plaques in patients with ACS in a study utilizing IVUS imaging. In CCTA, spotty calcification is defined as a small, dense (>130 HU) plaque component surrounded by noncalcified plaque tissue. The typical cut-off to define...
a small calcification in CCTA as spotty is <3 mm.\textsuperscript{16,36,53} Spotty calcifications have been further differentiated into small (<1 mm), intermediate (1–3 mm), and large (>3 mm) calcifications.\textsuperscript{81} Small spotty calcification has the strongest association with vulnerable plaque features defined by VH-IVUS.\textsuperscript{84} Furthermore, in multiple cross-sectional studies in patients with ACS and SAP, spotty calcification is associated with ACS culprit lesions.\textsuperscript{34,35–36} However, results vary widely, and highlight the current uncertainty in the relationship between spotty calcification and plaque rupture.\textsuperscript{72} With further improvements in CT technology, detection of microcalcifications, which have been suggested to be a frequent feature in unstable angina, might be feasible.\textsuperscript{82}

**Functional plaque characteristics**

Plaques develop at specific areas of coronary arteries where flow is disturbed, such as the outer walls of bifurcations, in side branches, and in the inner curve of arteries,\textsuperscript{83–87} despite risk factors for plaque formation (including smoking, high cholesterol levels, hypertension, and insulin resistance) affecting the whole vascular bed.\textsuperscript{83–87} Haemodynamic factors, such as endothelial shear stress (ESS), are pathologically important for the spatial localization and development of atherosclerotic plaques.\textsuperscript{88} Low ESS promotes an atherogenic milieu and high-risk plaque formation, whereas high ESS at stenotic vulnerable plaque sites promotes plaque rupture by destabilization of the fibrous cap.\textsuperscript{89–91}

In the early 1990s, *post-mortem* studies indicated that more than two-thirds of infarctions evolve from non-obstructive lesions (that is lesions occupying <70% of the lumen).\textsuperscript{92} However, histopathological investigations have now challenged these studies, and a high portion of culprit lesions now seem to cause obstructive luminal narrowing (>75% area stenosis was seen in 70% of plaque ruptures), especially in late stages of plaque development before the disruption of the fibrous cap.\textsuperscript{14,15,93} These observations correlate with evidence that patients with ischaemic lesions have a poor prognosis.\textsuperscript{84,95} Indeed, increased plaque vulnerability might in part be a consequence of haemodynamic perturbations and altered shear stress owing to abnormal fractional flow reserve (FFR).\textsuperscript{96} Invasive FFR is the gold standard method for the identification of lesions that result in ischaemia,\textsuperscript{97} and the combination of ESS and FFR might, therefore, provide a novel functional dimension in plaque vulnerability assessment (Figure 1). Advances in computational fluid dynamics (CFD) have enabled the simulation of coronary flow and pressure-based metrics on the 3D geometry of the coronary artery tree.\textsuperscript{98} When CFD is added to standardly acquired CCTA dataset, ESS-CT and FFR-CT coronary maps can be calculated.\textsuperscript{99–101}

**ESS-CT**

ESS is the tangential force generated by the friction of flowing blood on the endothelial surface of the arterial wall.\textsuperscript{102} In coronary artery segments with low and disturbed or turbulent flow—where ESS is low—the endothelial cell gene expression initiates a proatherogenic pattern.\textsuperscript{103–104} Persistently low ESS reduces nitric oxide production, increases LDL uptake, promotes endothelial cell apoptosis, and induces local oxidative stress and inflammation, which induce an atherogenic endothelial phenotype and subsequently leads to the development of high-risk lesions.\textsuperscript{86,88} By contrast, in straight arterial segments with undisturbed laminar flow—where ESS varies within a physiological range—endothelial cells express atheroprotective genes leading to plaque stability and quiescence.\textsuperscript{87,88,102} However, high shear stress at the stenotic portion of the plaque might initiate pathophysiological processes that promote plaque destabilization and rupture.\textsuperscript{87,89}

In serial IVUS studies of coronary arteries in diabetic pigs, the majority of vulnerable plaques developed in vessel segments characterized by persistently low ESS.\textsuperscript{55,88,105} Furthermore, the magnitude of low ESS at baseline was significantly associated with the severity of high-risk plaque features at follow-up.\textsuperscript{88} Another animal study has refined the concept that low ESS promotes coronary plaque growth and vulnerability by demonstrating that dyslipidaemia and low ESS have a synergistic effect leading to the development of vulnerable atheromas.\textsuperscript{106}

The first natural-history VH-IVUS study in humans assessed the left anterior descending artery in 20 patients with nonobstructive CAD at enrolment and at 6 months follow-up.\textsuperscript{90} Low ESS segments developed increased plaque area and necrotic core as well as constrictive remodelling, whereas high ESS segments developed greater necrotic core and regression of fibrous and fibrofatty tissue, and excessive positive remodelling, suggestive of transformation to a more-vulnerable phenotype.\textsuperscript{90} These observations highlight the importance of low ESS in vulnerable plaque development and high ESS in the
Plaques that rupture cause substantial luminal narrowing at the time of the acute event. Histopathological investigations demonstrated that plaques that rupture but are nonstenotic are very rare. The assessment of luminal narrowing at the site of a large lipid-rich plaque might, therefore, be an important addition to high-risk plaque features and could aid the identification of vulnerable plaques.

In a histopathological study of ruptured plaques and TCFAs, 70% produced significant narrowing (>75%) of the cross-sectional luminal area. The remaining 30% of nonobstructive ruptured plaques were further subdivided into those with luminal narrowing of 50–75% and those with luminal narrowing <50% (25% and 5% of lesions respectively). Importantly, the investigators assessed the nonruptured TCFAs, which are the potential targets for noninvasive imaging, and found 40% also caused luminal narrowing of >75%. Because these lesions are likely to cause angina, they are probably more likely to be treated. However, lesions with an intermediate stenosis can be large, but not necessarily associated with symptoms. Vulnerable plaques with a stenosis range of 50–75% (~50% of TCFAs) are the real targets for noninvasive imaging. Abbreviations: FFR, fractional flow reserve; TCA, thin cap fibroatheroma.

**Figure 6** In histopathological studies of patients who suffered sudden cardiac death, 40% of nonruptured TCFAs also caused >75% luminal narrowing. These TCFAs with significant luminal narrowing (>75%), are likely to cause angina, and therefore be treated. Lesions with an intermediate stenosis can be of danger, as they are large, but are not necessarily associated with symptoms. Vulnerable plaques with a stenosis range of 50–75% (~50% of TCFAs) are the real targets for noninvasive imaging. Abbreviations: FFR, fractional flow reserve; TCA, thin cap fibroatheroma.

destabilization of these plaques. In the Prediction of Progression of Coronary Artery Disease and Clinical Outcome Using Vascular Profiling of Shear Stress and Wall Morphology (PREDICTION) trial, a total of 506 patients underwent three-vessel IVUS examination and were assessed again at 1-year follow-up. The results demonstrated that large plaque burden and low ESS can independently predict plaques that progressively enlarge and develop substantial lumen narrowing.

3D coronary geometry visualization by CCTA enables CFD to be applied to ESS-CT calculations and subsequent coronary wall behaviour assessment (Figure 5). These observations have been confirmed in a study using CCTA and IVUS for vascular profiling. CCTA was sufficiently accurate to determine ESS distribution in the main vessels and in the bifurcation regions. The CFD simulations in CCTA can be used to remove all plaques in a virtual environment to replicate the healthy vascular wall before the development of atherosclerotic plaques. In an exploratory investigation, static and dynamic parameters of ESS-CT were calculated in a virtual healthy coronary lumen to determine the best haemodynamic predictor of future plaque location. The results of this virtual experiment suggested that low ESS is a prerequisite for plaque formation; however, its presence alone is insufficient to predict future plaque locations. Dynamic factors that describe the time-dependent directional changes in ESS might, therefore, have an incremental prognostic value regarding plaque progression and vulnerability.

**FFR-CT**

FFR-CT will help in the identification of lesions with ischaemia and likely improve CT accuracy for the detection of high-risk lesions. Importantly, FFR-CT can be derived from CCTA, without the need for additional imaging, extra radiation, or any medication (Figure 7). Furthermore, FFR-CT provides a comprehensive three-vessel FFR from a single CCTA test, enabling FFR readings at any location of the coronary tree. Two prospective clinical trials have demonstrated that FFR-CT...
Importantly, the definition of vulnerable plaques by plaque assessment with CCTA. The theoretical volume-rendered CT image consists of FFR-CT with myocardial perfusion imaging (pink indicates normally perfused myocardium, light purple represents moderate ischaemia, whereas purple indicates severe perfusion defect), which can differentiate between epicardial and small vessel disease. In this specific case epicardial disease can be confirmed as FFR-CT and the perfusion map show similar ischaemic regions. Each coronary plaque receives a vulnerability score based on morphological and functional data. In this vulnerability score, 1 represents a stable lesion (green) and 10 a high-risk, rupture prone plaque (red). The figure demonstrates that plaques with no haemodynamic significance might also contain high-risk characteristics and have an increased vulnerability score. Abbreviations: CCTA, coronary computed tomography angiography; FFR, fractional flow reserve.

**Future directions for CCTA**

Coupling individual plaque morphology with plaque-specific functional data, including ESS and FFR, provide new opportunities for the noninvasive detection of plaques vulnerable to rupture. Combining morphological and functional metrics of coronary plaques (comprehensive plaque assessment) might lead to the development of a ‘vulnerable plaque score’ to indicate potential acute clinical events (Figure 8). Moreover, the high sensitivity and specificity of CCTA to identify atherosclerotic plaques can be combined with whole genome sequencing and epigenomics, whole transcriptome sequencing, unbiased proteomics, metabolomics, lipidomics, and lipoprotein proteomics to enable a stratified, step-wise approach (‘omics’ followed by CCTA if needed) to identify patients with subclinical CAD and vulnerable plaques, as in the Genetic Loci and the Burden of Atherosclerotic Lesions Study, for example. Combining the complex information obtained from CT with bioinformatics analysis of genomic or proteomic data, as well as data derived from electronic medical records and drug information, might realize a personalized approach to cardiovascular disease prevention and care for each patient.

Automated plaque assessment tools and computation fluid dynamics simulations will be necessary to provide multidimensional information on individual plaques. Importantly, the definition of vulnerable plaques by
CCTA has not yet been established. Large, longitudinal imaging trials to test the prognostic value of comprehensive plaque assessment and derive standardized CT metrics of vulnerable plaques are also warranted. Using hybrid scanners and novel contrast agents to acquire metabolic information might also improve CT plaque assessment. For example, inflammatory immune cell infiltration into the fibrous cap is a strong marker of plaque rupture vulnerability. In rabbits, the detection of inflammatory cells in atherosclerotic plaques by CCTA with iodinated nanoparticle contrast agent N1117 has been demonstrated. Furthermore, spectral CT imaging of gold-labelled HDL nanoparticles targeted to activated macrophages showed promising results in a mouse model of atherosclerosis. However, neither technique has been adapted for human patients. The high metabolic activity of macrophages and their metabolic dependence on exogenous glucose provide opportunity for combined positron emission tomography and CT imaging using a radiolabelled glucose analogue—fluorine-18-fluorodeoxyglucose (18F-FDG)—to noninvasively detect vulnerable, inflamed plaques. In a clinical investigation ‘culprit lesion site of patients with recent ACS as compared to stented lesions in patients with SAP. In a prospective clinical trial, 18F-sodium fluoride (18F-NaF) uptake co-localized to ruptured plaques in patients with ACS, and in patients with SAP 18F-NaF identified plaques with high-risk features, as determined by IVUS. The intriguing results showed that 18F-NaF positron emission tomography–CT imaging can detect metabolically active plaques by identifying areas of ongoing calcification activity.

Conclusion

CCTA has the ability to depict all main epicardial coronary branches and, therefore, enables both individual plaques and global coronary plaque burden to be evaluated. Modern CCTA assessment can identify multiple high-risk features—such as NRS, positive remodelling, low CT attenuation, and low ESS—that make a plaque vulnerable to rupture and, therefore, put individuals at increased risk of an acute cardiovascular event. Conversely, the limited spatial and contrast resolutions of CT scanners prevent the detection of some histological features of vulnerable plaques, such as fibrous cap thickness or plaque rupture. Nanoparticle contrast agents and hybrid imaging provide valuable information regarding the metabolic activity of atherosclerotic plaques, but initial results, although encouraging, need to be confirmed in large prospective trials. Patients with acute and stable chest pain syndromes—and perhaps even asymptomatic patients—might receive improved, targeted therapy and derive the health benefits from such interventions. However, CCTA uses ionizing radiation and its current use must, therefore, be limited to symptomatic patients with established clinical indication for this noninvasive test.

Generalized, large, prospective trials are needed to confirm the results of small, prospective or registry-based studies, because current data suggest that CCTA-based assessment of both individual and global coronary plaque features is highly predictive of future adverse events. Randomized trials testing whether interventions made on the basis of CCTA assessment can improve both health and economic outcomes should also be conducted. Further strategies targeting vulnerable plaques, or to treat patients more effectively, might also include medical therapy and/or plaque sealing with a biodegradable vascular scaffold.

Review criteria

English-language articles published between January 2000 and 2014 were identified by searching the PubMed database. The search criteria included were: “coronary ct angiography” or “non-invasive imaging”, alone and in combination with “vulnerable plaque”, “coronary plaque”, “atherosclerotic plaque”, “high-risk plaque”, “computational fluid dynamics” and “functional assessment”. Additional articles were identified from the reference lists of retrieved articles.


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Author contributions

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